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(54) Title: POURABLE FRYING COMPOSITION

(57) **Abstract:** Water continuous, pourable compositions comprising more than 50 to 80 wt % fat, an emulsifier having a hydrophilic/lipophilic balance value of at least 7, an antisputtering composition and optionally a biopolymer in an amount of at most at most 0.3 wt % on total composition weight are suitable frying compositions.

Pourable frying composition

**Field of the invention**

5 The invention relates to pourable, water continuous frying compositions which are storage stable and show good spattering behaviour and frying characteristics.

**Background to the invention**

10

There is a continuous desire among consumers for frying products which are easily dosed and do not show spattering upon use as shallow frying agent.

15 Spattering of a common frying medium, such as margarine which is a water in oil emulsion, is believed to be caused by superheating of water droplets. At a certain point after heating the frying medium said water droplets explosively evaporate, whereby oil can be spread all over the surroundings  
20 of a frying pan wherein the emulsion is heated. This may cause danger to the person who intends to fry foodstuff in the heated emulsion.

Another disadvantage of frying products that is often encountered is residue formation. Brown or black residue may be  
25 formed when for example biopolymers such as proteins are heated in a frying pan. Preferred products show little or no residue formation when used as a frying agent.

Common frying agents such as butter or margarine are fat  
30 continuous and hence show the undesired spattering behaviour as explained above.

Due to their easy dosing, liquid margarines have gained popularity with consumers.

US-A-4,292,333 relates to low fat content butter flavoured 5 liquid spreads which simulate the flavour, texture, mouthfeel, appearance and stability of commercial liquid margarine.

US-A- 4,273,790 discloses a low fat liquid spread comprising less than 40 wt% fat, a continuous aqueous phase comprising a 10 stabiliser and an emulsifier system comprising a combination of a lipophilic emulsifier and a hydrophilic emulsifier, thus providing a stable liquid emulsion at 40 °F.

GB-A-1,359,639 discloses pourable margarines having increased 15 stability against oil and water separation which are suitable for shallow frying.

However none of the above documents provides a frying product which is easily pourable, shows good spattering behaviour, low 20 sediment on shallow frying use and a stability against serum separation on shelf. Therefore it is an object of the invention to provide products that show these characteristics.

According to a further object the products are heat stable such 25 that they can be subjected to sterilisation or pasteurisation treatments.

#### **Summary of the invention**

30 It has now surprisingly been found that a composition which comprises a specified emulsifier, a specific fat content dispersed in an aqueous phase and an antispattering agent, shows very good spattering behaviour, is storage stable and

does not lead to a lot of burnt sediment formation upon use as a frying agent.

Therefore the invention relates to a pourable, water continuous 5 frying composition having a Bostwick value at 15 °C of at least 5, comprising 40 to 80 wt% fat, an antispattering agent, at least one emulsifier having a hydrophilic/lipophilic balance value of at least 7, and optionally a biopolymer in an amount of at most 0.3 wt% on total composition weight.

10

In a second aspect the invention relates to a process for the preparation of this emulsion.

15 In a further aspect the invention relates to use of this emulsion for shallow frying.

#### **Detailed description of the invention**

Products according to the invention are products with a 20 Bostwick value of at least 5 at 15 °C. The Bostwick value is measured as described in the examples.

Preferably the Bostwick value for products according to the invention is 8 - 24, more preferably 11 - 21, most preferably 15 - 18 at 15 °C.

25

Spattering can be measured by determining the spattering value according to the method illustrated in the examples. Preferably food products according to the invention show a primary spattering value, SV1, (spattering upon heating of a frying 30 product such as margarine, without incorporation of a food product to be fried) of from 7 to 10, more preferably from 8.5 to 10. The secondary spattering value, SV2, (spattering upon incorporation of a food product such as meat in a shallow

frying product) for products according to the invention is preferably from 5-10.

Upon use of the composition according to the invention as a  
5 frying agent, unexpected good primary and secondary spattering behaviour was observed. Furthermore, despite the relatively low fat content, the emulsions retained good pourability compared to fat continuous pourable frying compositions in the same fat range. Compared to the known fat continuous liquid margarines  
10 of varying fat content, the current compositions show better pourability at the same fat content which results in easier dosing, a non fatty appearance and less residue in a bottle from which the composition is poured. Furthermore the current compositions do not require the presence of a hardstock fat  
15 which can be advantageous.

Moreover the claimed compositions are storage stable.

Storage stability is defined as having a water separation layer at the bottom of a glass jar of below 7 vol% after storage for  
20 2 weeks at 15 °C. Preferably the water separation layer is below 5 vol%, more preferred below 2 vol% of the total product volume.

Advantageously this stability can be obtained without the need for a solid fat being present for structuring.

25

Heat stability is defined as an oil phase separation of less than 3 vol% separated oil after storage of the composition in a glass container at 121 °C for 10 minutes. The separation volume is measured directly after the heat treatment.

30

The compositions according to the invention are water continuous products which comprise an aqueous phase and a dispersed fat phase.

In order to ensure that the emulsion of the fat in the aqueous 5 phase is stable, the products comprise at least one emulsifier which is characterised by a hydrophilic/lipophilic balance value of at least 7. This value is abbreviated as HLB value. The HLB value is a well known measure of balance between the hydrophilicity and the lipophilicity of an emulsifier. For a 10 discussion of this value reference is made to "An introduction to food colloids" by Eric Dickinson, Oxford University Press, 1992, page 47-49.

Preferably the emulsifier is selected from the group comprising 15 di-acetyl tartaric acid esters of monoglycerides and/or diglycerides (DATEM), polyoxyethylene sorbitan fatty acid esters (Tween), sucrose esters, sodium stearoyl lactylate (SSL), polyglycerol esters (PGE), acetylated pectin, esters of citric acid with monoglycerides and/or with diglycerides (CAE), 20 lactic acid esters of mono-and/or diglycerides, succinic acid esters of mono-and/or diglycerides; or combinations thereof.

More preferably the emulsifier is selected from the group: di-acetyl tartaric acid esters of monoglycerides and/or 25 diglycerides (DATEM), polyoxyethylene sorbitan fatty acid esters (Tween), sucrose esters, polyglycerol esters (PGE) and esters of citric acid with monoglycerides and/or with diglycerides (CAE).

30 Di-acetyl tartaric acid esters of mono-and/or diglycerides (DATEM), were found to lead to highly stable emulsions which showed a primary spattering value of at least 8 and secondary

spattering value of at least 5 and often even at least 7 and giving no sediment after shallow frying.

Therefore in a preferred embodiment, the emulsifier is a di-acetyl tartaric acid ester of mono-and/or diglycerides (DATEM).

5

Sucrose ester were found to lead to highly stable emulsions which showed a primary spattering value of at least 8 and secondary spattering value of at least 5 and often even at least 7 and giving no sediment after shallow frying.

10 Although single emulsifiers are preferred in view of costs and ease of processing, a combination of emulsifiers fulfilling the HLB value requirement is also encompassed in the invention. Besides the emulsifier with HLB value of at least 7 other emulsifiers can be present, but their presence is not required

15 to fulfil at least some of the objects of the current invention.

The amount of emulsifier in the claimed frying composition is such that the emulsion is storage stable, heat stable and still

20 shows good frying behaviour.

It will be appreciated that the amount of emulsifier added varies with the type of emulsifier and the ratio between the fat phase and the aqueous phase of the emulsion.

Generally, the lower the amount of fat, the lower the amount of

25 emulsifier can be.

Suitable amounts of emulsifier were found to be between 0.1 and 5 wt% on total frying composition weight. Preferably the total amount of emulsifier is from 0.35 to 5 wt%, more preferably

30 from 0.5 to 3 wt%.

In a highly preferred embodiment, the invention relates to a pourable water continuous frying composition having a Bostwick value at 15 °C of at least 5, comprising 40 to 80 wt% fat, and 0.3 to 3 wt%, preferably 0.35 to 3 wt% of di-acetyl tartaric acid ester of mono- and/or diglycerides.

In addition to an emulsifier, the composition comprises an antispattering agent to further improve frying characteristics.

Any suitable antispattering agent can be used, but it was found 10 that a combination of salt and a lecithin leads to particularly good results.

Although these compounds are well known antispattering agents for fat continuous frying agents, their positive effect in a pourable, water continuous composition is unexpected.

15

Therefore in a preferred embodiment, the antispattering composition comprises salt in an amount of from 0.1 to 5 wt%, preferably 0.1 to 3 wt% on total weight of the frying composition and a lecithin in an amount of from 0.05 to 2 wt%, 20 preferably 0.05 to 1 wt% on total weight of the frying composition.

The total amount of antispattering agent is preferably from 0.15 to 7 wt% on total composition weight.

25

Lecithin can be obtained in the form of several types, depending on the process used for its preparation.

For the purpose of the invention, lecithins, which comprise phosphoacylglycerols, can be divided in three groups according 30 to their preparation. The first group is formed by native lecithins such as Bolec ZT<sup>TM</sup>.

Native lecithins are for example obtained from triglyceride oils which have been filtered, extracted and stripped.

The second group of lecithins is formed by (partly) hydrolysed 5 lecithins which originate from native lecithins which have been hydrolysed for example by use of the enzyme phospholipase A or by chemical hydrolysis. Hydrolysed lecithins can also be prepared by chemical synthesis.

A third group of lecithins comprises fractionated lecithins 10 such as the alcohol soluble fraction of native lecithins such as Cetinol<sup>TM</sup>.

This type of lecithin can be obtained in a process wherein native lecithins are extracted with alcohol.

15 Examples of these three groups of lecithins are:

native lecithin: Bolec ZT<sup>(tm)</sup>, Adlec<sup>(tm)</sup>, Sternpur PM<sup>(tm)</sup>;

hydrolyzed lecithin BOLEC MT<sup>(tm)</sup>, Sternphil<sup>(tm)</sup>, Adlec E<sup>(tm)</sup>;

fractionated lecithin: Cetinol<sup>(tm)</sup>, Nathin 3-KE<sup>(tm)</sup>.

20 For the purpose of the invention lecithins are from vegetable origin. Lecithins in each group can optionally be de-oiled lecithins.

In a preferred embodiment lecithin is fractionated lecithin.

25 The amount of fractionated lecithin is preferably 0.15 - 3 wt%, more preferably 0.2 - 2wt%, most preferably 0.2 - 1wt%.

Alternative antispattering agents include dispersed gas phase, sand, hydrophobic particles such as silica particles, citric acid esters.

30

Optionally the composition according to the invention comprises a biopolymer to improve emulsion stability. More preferred this biopolymer is selected from the group comprising proteins,

starches, pectins, carrageenans, alginates, galactomannans (e.g. guar gum, locust bean gum), celluloses and modified celluloses, bacterial exopolysaccharides (e.g. xanthan, gellan).

5

The amount of biopolymer when added is below 0.3 wt% to ensure that excessive sediment formation and burning are prohibited.

Most preferred, biopolymer is present in an amount of from 0.01  
10 to 0.3 wt%.

The pH of the composition according to the invention is preferably from 3 to 8.

For reasons of microbiological stability the pH can be lowered  
15 to between 0.5 and 5, preferably 2.5 - 5, by the use of any suitable, food grade acid.

The composition according to the invention comprises a fat in an amount of from 40 to 80 wt%.

20 Preferred compositions comprise more than 50 to 80 wt% fat, more preferably 51 - 75 wt% fat; even more preferably 55 - 75 wt% fat, most preferably 55 - 65 wt% fat.

The fat can be any fat, but a fat rich in triglycerides  
25 comprising (poly) unsaturated fatty acid residues is highly preferred. As mentioned above one of the advantages of the claimed composition is that solid fat (such as hardened rapeseed oil) is not required to obtain a stable composition.

For common, fat continuous, pourable frying compositions, a  
30 solid fat is usually added to improve the product stability. These solid, hardened fats contain high amounts of saturated fatty acids, which are generally considered less beneficial for

health than fats rich in triglycerides comprising (poly) unsaturated fatty acid residues.

Therefore the fat is preferably selected from the group  
5 comprising sunflower oil, soybean oil, rapeseed oil, peanut oil, safflower oil, cottonseed oil, olive oil, corn oil, groundnut oil, or low melting butterfat fractions and/or combinations thereof. These fats may be partially hydrogenated.

10 Most advantageously the fat or combination of fats is selected such that the solid fat content of the fat or fat blend is 0% at 15 °C and above.

The fatty phase can also comprise sucrose polyesters (SPE's).

15

The composition is water continuous which implies that the fat is dispersed in a continuous aqueous phase in the form of fat droplets.

Storage stable compositions which show little creaming  
20 advantageously are characterised by an average fat droplet size ( $d_{43}$ ) of the fat of less than 8 µm, preferably less than 6 µm, more preferred from 0.35 to 4 µm.

In addition to the above-mentioned ingredients, compositions  
25 according to the invention may optionally contain further ingredients suitable for use in these products. Examples of these materials are sugar or other sweetener materials, EDTA, spices, salt (other than antispattering salts), bulking agents, egg yolk, anti-oxidants, flavouring materials, colouring  
30 materials, acids, preserving agents, and fruit and/or vegetable particles.

In a further aspect the invention relates to a process for the preparation of the claimed frying composition.

In general any suitable process can be used.

- 5 Preferably the process comprises the steps of emulsification of a fat phase comprising fat phase ingredients with an aqueous phase comprising aqueous phase ingredients such that the resulting average fat droplet size  $d_{43}$  is below 8  $\mu\text{m}$ .
- 10 Fat phase ingredients are those ingredients which are either fat soluble or fat dispersible. Examples are fat soluble emulsifiers and antispattering agents such as lecithin, oil soluble flavour/colouring composition, colourants, vitamins and anti-oxidants.

15 Aqueous phase ingredients are those ingredients which are either water soluble or water dispersible. Examples of such ingredients are stabilisers and protein, salt, preservatives, acidifiers, emulsifiers.

20 Biopolymer, when added, is preferably added to the aqueous phase before it is mixed with the fat phase.

In a preferred embodiment, the aqueous phase comprising aqueous

- 25 phase ingredients is heated to at least 50 °C before mixing with the oil phase.

Preferably the fat phase is added to the aqueous phase slowly, while mixing.

- 30 Emulsification by use of a homogeniser, a colloid mill or a high shear mixer or similar apparatus is preferred.

If the emulsifier is a di-acetyltauric acid ester of mono- and/or diglycerides the aqueous phase is preferably set to pH of 4 or higher and subsequently emulsified with a fat phase.

5 Alternatively if polyglycerol esters are the emulsifiers used, the pH of the composition is between 5 and 8 for reasons of increased stability of the composition under these conditions.

In a further aspect the invention relates to the use of the  
10 composition according to the invention for shallow frying of foodstuff. During shallow frying a volume of composition is heated in a frying pan to about 160 to 200 °C. Once the frying medium has reached the desired temperature, food stuff to be fried is put into the hot medium.

15 Examples of foodstuff which can suitably be fried with the composition according to the invention include meat, vegetables, eggs, fish.

The invention will be illustrated by the following examples.

20

### **Examples**

#### Analysis methods

25 **Creaming**

Emulsions are tested for stability visually at 15 °C. Emulsion is filled into 100 ml measuring cylinders (diameter about 3.5 cm). The amount of free water that is visible at the bottom of the cylinder is measured after 2 weeks in terms of a volume  
30 percentage on total emulsion volume.

**Droplet size**

Droplet size distribution of the emulsion is measured by small angle laser light scattering using a Malvern Mastersizer. The average droplet size ( $d_{43}$ ) of the fresh emulsion is the  
5 parameter of interest.

**Bostwick value determination**

Pourability is measured according to the standard Bostwick protocol. The Bostwick equipment consists of a 100 ml reservoir  
10 with an outlet near the bottom of a horizontally placed rectangular tub and closed with a vertical barrier.

The tub's bottom is provided with a 24 cm measuring scale, extending from the outlet of the reservoir. When equipment and sample both have a temperature of 15 deg. C, the reservoir is  
15 filled with 100 ml of the sample after it has been shaken by hand ten times up and down. When the closure of the reservoir is removed the sample flows from the reservoir and spreads over the tub bottom.

The path length of the flow is measured after 15 seconds. The  
20 value, expressed as cm per 15 seconds is the Bostwick rating, which is used as measure of pourability.

**Spattering value**

The spattering behaviour of compositions according to the  
25 invention was evaluated after storage of the products 14 days at 15 °C.

Primary spattering was assessed under standardised conditions in which an aliquot of the composition was heated in a glass dish and the amount of fat spattered onto a sheet of paper held  
30 at a fixed distance above the dish was assessed after the water content of the food product had been driven off by heating.

In assessment of the primary spattering value about 35 g of the composition was heated in a glass dish on an electric plate set at 205 °C. (The amount of composition heated depends on the fat content. If the composition contains 60% fat, about 35 g is used, if the composition contains more fat, less product is heated arriving at about 21 g to be heated for a product containing about 97% fat.) The fat that spattered out of the pan by force of expanding evaporating water droplets was deposited on a sheet of paper situated above the pan. The image obtained was compared with a set of standard pictures number 0-10 whereby the number of the picture which most closely resembled the product was recorded as the spattering value for that product. A value of 10 indicates no spattering (no fat loss) and zero indicates very bad spattering (almost total fat loss). The general indication is as follows.

<i>Score</i>	<i>Comments</i>
10	Excellent
8	Good
5	Acceptable
4	Unsatisfactory
2	Very Poor

#### **Example 1A**

##### **20 Product composition**

<i>Ingredient</i>	<i>Amount in wt%</i>
-------------------	----------------------

Sunflower Oil	59 %
Water	balance to 100wt%
Datem 1935	1 %
Salt (NaCl)	1 %

15

Lecithin (cetinol)	1 %
Potassium sorbate	0.1 %

pH 5.2

5

DATEM is di-acetyl tartaric acid ester of mono-and diglycerides obtainable from Quest.

**Example 1B**

10

Product composition was the same as for example 1A with the modification that example 1B additionally contained 0.1 wt% xanthan gum. Furthermore the pH of the aqueous phase of this example was set to 4.2.

15

**Example 2***Product composition*

Ingredient	Amount in wt%
Sunflower Oil	59%
Water	balance to 100 wt%
Datem 1935	1%
Salt (NaCl)	1.5%
25 Lecithin (cetinol)	1 %
Potassium sorbate	0.1%
Guar gum	0.1%

pH 4.3

30

*Process for example 1A, example 1B and example 2*

Water continuous emulsions were prepared starting with the water phase in which the o/w emulsifier and other water-soluble ingredients were dissolved. The pH of the water phase was adjusted to 5.2 (ex 1A), resp. 4.2 (ex 1B), resp. 4.3 (ex 2).

5 The oil phase, in which lecithin was dissolved was slowly added to the water phase after which the two phases were mixed for 5 minutes at medium speed using an ultra-turrax (Janke & Kunkel). The temperature during the emulsification was 40 to 45°C. The emulsions were poured into a 100-ml measuring cylinder and in a  
10 glass jar. They were stored at 15°C.

### **Example 3**

#### *Product composition*

15	Ingredient	Amount in wt%
	Sunflower Oil	60 %
	Water	balance to 100 wt%
	Sucrose ester(S-1570)	1 %
20	Salt (NaCl)	1.5 %
	Lecithin (cetinol)	0.3 %
	Potassium sorbate	0.1 %

pH 4 - 4.5

25

### **Example 4**

#### *Product composition*

30	Ingredient	Amount in wt%
	Sunflower Oil	60 %
	Water	balance to 100wt%

Polyglycerol ester	
(Triodan)	1 %
Salt (NaCl)	1.5 %
Lecithin (cetinol)	0.3 %
5 Potassium sorbate	0.1 %

**Example 5***Product composition*

10 Ingredient	Amount in wt%
Sunflower Oil	60 %
Water	balance to 100wt%
Citric acid ester (Citrem N12)	1 %
15 Potassium sorbate	0.1 %

**Example 6***Product composition*

20 Ingredient	Amount in wt%
Sunflower Oil	60 %
Water	balance to 100wt%
Polyoxyethylene sorbitan	
25 fatty acid ester (Tween 60)	1 %
Salt (NaCl)	1.5 %
Lecithin (Cetinol)	0.3 %
Potassium sorbate	0.1 %

**Example 7***Product composition*

Ingredient	Amount in wt%
5 Sunflower Oil	50.5 %
Water	balance to 100 wt%
Datem 1935	1.2%
Salt (NaCl)	1.5 %
Lecithin (cetinol)	0.3 %
10 Potassium sorbate	0.1 %

pH 4.5

*Process for examples 3 to 7*

15 Water for the aqueous phase was heated to around 75 °C, and the emulsifier was added while mixing. For example 3 the emulsifier was slurried with a small amount of cold water before addition to hot water. After mixing with a Silverson mixer to disperse the emulsifier, the other water soluble ingredients were added to the aqueous phase. The pH was adjusted to the desired value if required.

The oil phase ingredients (which include lecithin) were mixed and heated to around 60 °C. The oil phase was then added slowly to the aqueous phase while mixing. After addition of

20 the oil phase was complete the mixing speed was turned up to high and mixing continued for 5 minutes more. The premix was then homogenized by recirculating for 3 minutes through a high pressure homogenizer operating at 200 bar.

25

The resulting products for examples 1-7 were scored as follows:

Measure ment	Ex. 1A	Ex. 1B	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6	Ex. 7
Creaming	2%	0	0	Trace	3%	2%	Trace	0
Bostwick value (cm/15 sec)	20	16	8	9	13	24	24	24
average droplet size ( $d_{43}$ )	n.d.	n.d.	n.d.	2.7	2.0	3.7	1.3	1.1
Frying behaviour								
Sediment formation	none	none	None	None	none	none	None	None
Shallow frying SV1 / SV2	8 / 6	7 / 5	8 / 6.5	9 / 7	7 / 8	9 / 3	7 / 5	9 / 7

5

n.d.: not determined

### Example 8

#### 10 Product composition

Ingredient	Amount in wt%
------------	---------------

Sunflower Oil 59.65 %

Water balance to 100wt%

#### 15 Datem 1935

Salt (NaCl) 1.5 %

Lecithin (cetinol) 0.3 %

Potassium sorbate 0.1 %

20

Guar gum	0.1 %
Sodium hydroxide solution (20wt%)	0.15 %
Beta carotene	0.05 %

5 300 kg emulsion of the above composition was prepared. Ingredients were premixed in a low shear mixer and after pasteurisation (80 °C) homogenized in a high pressure homogenizer operating at 200 bar. The resulting emulsion was cooled to 15 °C using a tubular heat exchanger.

10

The emulsion stability of the resulting emulsion was determined. At 15 °C and 25 °C the emulsion was stable for at least two months. At 30 °C it was stable for at least 6 weeks and at 35 °C at least 5 weeks. The spattering performance was:

15 SV1: 7.5 and SV2: 6.0. The Bostwick value was 14 cm/15 seconds.

**Claims**

1. Pourable, water continuous frying composition having a Bostwick value at 15 °C of at least 5, comprising more than 50 to 80 wt% fat, an antispattering agent, at least one emulsifier having a hydrophilic/lipophilic balance value of at least 7, and optionally a biopolymer in an amount of at most 0.3 wt% on total composition weight.
2. Pourable composition according to claim 1 wherein the antispattering agent comprises salt in an amount of from 0.1 to 5 wt% on total weight of the frying composition and a lecithin in an amount of from 0.05 to 2 wt% on total weight of the frying composition.
3. Pourable composition according to claim 1 or 2 wherein the emulsifier is selected from the group comprising di-acetyl tartaric acid esters of monoglycerides and/or diglycerides (DATEM), polyoxyethylene sorbitan fatty acid esters (Tween), sucrose esters, sodium stearoyl lactylate (SSL), polyglycerol esters (PGE), acetylated pectin, esters of citric acid with monoglycerides and/or with diglycerides, lactic acid esters of mono-and/or diglycerides, succinic acid esters of mono-and/or diglycerides; or combinations thereof.
4. Pourable composition according to any of claims 1-3 comprising 0.1 to 5 wt% of emulsifier.
5. Pourable composition according to any of claims 1-4 wherein the emulsifier is DATEM in a preferred amount of from 0.3 to 3 wt%.

6. Pourable composition according to any of claims 1-5 characterised by a pH of between 3 and 8.
7. Pourable composition according to any of claims 1-6 comprising a biopolymer.
8. Pourable composition according to claim 7 wherein the biopolymer is present in an amount of from 0.01 to 0.3 wt%.
9. Pourable composition according to any of claims 1-8 wherein the fat is dispersed in a water phase, whereby the average droplet size ( $d_{43}$ ) of the fat is less than 8  $\mu\text{m}$ , preferably less than 6  $\mu\text{m}$ , more preferred from 0.35 to 4  $\mu\text{m}$ .
10. Process for the preparation of a pourable, water continuous frying composition according to any of the previous claims, comprising the steps of emulsification of a fat phase comprising fat phase ingredients with an aqueous phase comprising aqueous phase ingredients such that the resulting average fat droplet size  $d_{43}$  is below 8  $\mu\text{m}$ .
11. Process for the preparation of a pourable, water continuous frying composition according to claim 5 wherein an aqueous phase comprising a di-acetyltauric acid ester of mono- and/or diglycerides is set to a pH of 4 or higher and subsequently emulsified with a fat phase.
12. Use of the composition according to any of claims 1-9 for shallow frying of foodstuff.

## INTERNATIONAL SEARCH REPORT

In tional Application No  
PCT/EP 01/13310

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC 7 A23D7/00 A23D7/02

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A23D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

PAJ, EPO-Internal, WPI Data, FSTA

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	PATENT ABSTRACTS OF JAPAN vol. 1998, no. 10, 31 August 1998 (1998-08-31) -& JP 10 113145 A (NISSHIN OIL MILLS LTD:THE), 6 May 1998 (1998-05-06) abstract ---	1-12
Y	PATENT ABSTRACTS OF JAPAN vol. 1997, no. 10, 31 October 1997 (1997-10-31) -& JP 09 163952 A (NISSHIN OIL MILLS LTD:THE), 24 June 1997 (1997-06-24) abstract ---	1-12
Y	EP 0 716 811 A (NESTLE SA) 19 June 1996 (1996-06-19) the whole document ---	1-12 -/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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Date of the actual completion of the international search

Date of mailing of the international search report

26 February 2002

05/03/2002

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## INTERNATIONAL SEARCH REPORT

Int'l Application No  
PCT/EP 01/13310

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 253 429 A (UNILEVER NV ;UNILEVER PLC (GB)) 20 January 1988 (1988-01-20) column 7, line 24 - line 42 examples 1-4 claims 16-19 ----	1-12
A	WO 00 38546 A (UNILEVER PLC ;LEVER HINDUSTAN LTD (IN); BENJAMINS JAN (NL); UNILEV) 6 July 2000 (2000-07-06) page 5, line 4 - line 7 page 17, line 9 - line 27 ----	1
A	US 4 292 333 A (BOSCO PETER M ET AL) 29 September 1981 (1981-09-29) cited in the application the whole document ----	1
A	GB 1 359 639 A (UNILEVER LTD) 10 July 1974 (1974-07-10) cited in the application the whole document -----	1

## INTERNATIONAL SEARCH REPORT

Information on patent family members

Int'l Application No

PCT/EP 01/13310

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
JP 10113145	A	06-05-1998	NONE		
JP 09163952	A	24-06-1997	JP	2921664 B2	19-07-1999
EP 0716811	A	19-06-1996	EP	0716811 A1	19-06-1996
			AT	199484 T	15-03-2001
			BR	9505744 A	23-12-1997
			DE	69426823 D1	12-04-2001
			DE	69426823 T2	13-06-2001
			DK	716811 T3	09-07-2001
			ES	2155461 T3	16-05-2001
			NZ	280647 A	27-05-1998
			PL	311776 A1	24-06-1996
			PT	716811 T	31-07-2001
			US	5773072 A	30-06-1998
			ZA	9510571 A	12-06-1997
EP 0253429	A	20-01-1988	AT	64828 T	15-07-1991
			AU	600953 B2	30-08-1990
			AU	7479187 A	07-01-1988
			CA	1328052 A1	29-03-1994
			DE	3771137 D1	08-08-1991
			EP	0253429 A1	20-01-1988
			JP	1897954 C	23-01-1995
			JP	6026658 B	13-04-1994
			JP	63036827 A	17-02-1988
			US	5079028 A	07-01-1992
			ZA	8704722 A	22-02-1989
WO 0038546	A	06-07-2000	AU	2281000 A	31-07-2000
			AU	2281100 A	31-07-2000
			BR	9916447 A	04-09-2001
			BR	9916510 A	04-09-2001
			CZ	20012327 A3	14-11-2001
			CZ	20012328 A3	14-11-2001
			WO	0038546 A1	06-07-2000
			WO	0038547 A1	06-07-2000
			EP	1139799 A1	10-10-2001
			EP	1139800 A1	10-10-2001
US 4292333	A	29-09-1981	AU	538724 B2	23-08-1984
			AU	6451680 A	28-05-1981
			DE	3043656 A1	20-08-1981
			DK	492980 A	20-05-1981
			ES	496984 D0	01-05-1982
			ES	8204463 A1	01-08-1982
			FI	803590 A ,B,	20-05-1981
			FR	2469876 A1	29-05-1981
			GB	2066641 A ,B	15-07-1981
			IT	1134322 B	13-08-1986
			JP	56148240 A	17-11-1981
			MX	6753 E	25-06-1986
			NL	8006316 A	16-06-1981
			SE	450452 B	29-06-1987
			SE	8007997 A	20-05-1981
			WO	8101353 A1	28-05-1981
			US	4468408 A	28-08-1984

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 01/13310

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
GB 1359639	A 10-07-1974	AT 320410	B	10-02-1975
		AU 465031	B	18-09-1975
		AU 3553671	A	17-05-1973
		BE 775304	A1	12-05-1972
		CA 938493	A1	18-12-1973
		CH 555644	A	15-11-1974
		DE 2156121	A1	18-05-1972
		ES 396966	A1	01-03-1975
		FR 2113957	A5	30-06-1972
		IE 35802	B1	26-05-1976
		IT 1043878	B	29-02-1980
		JP 51016507	B	25-05-1976
		NL 7113682	A	11-04-1972
		NL 7115372	A	16-05-1972
		SE 381798	B	22-12-1975
		ZA 7107558	A	25-07-1973